Estrogen Receptor β Antibody



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Applications: W	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 52, 55, 63	Source/Isotype: Rabbit	UniProt ID: #Q92731	Entrez-Gene Id: 2100
Product Usage Information		Application Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Estrogen Receptor β Antibody detects endogenous levels of total Estrogen Receptor β protein. This antibody is predicted to cross-react with all Estrogen Receptor β isoforms. This antibody does not cross react with Estrogen Receptor α .				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human Estrogen Receptor β 1 protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Estrogen Receptor β (ER β) is a member of the nuclear receptor superfamily of transcription factors and was discovered to be encoded by a gene (<i>ESR2</i>) distinct from that encoding Estrogen Receptor α (ER α) (1,2). While studies have revealed that alternative splicing generates mutiple isoforms of ER β that differ at their respective C-termini and in tissue distribution, ER β 1 is believed to be the longest and only fully functional isoform (3,4). Indeed, it has been reported that shorter isoforms of ER β (ER β 2, β 4, and β 5) can heterodimerize with ER β 1 and enhance its transcriptional activity in an estradiol-dependent manner (4). ER β is expressed in a wide range of normal and malignant tissues, many of which coexpress ER α . It is proposed that ER β has an antiproliferative role, perhaps through heterodimerization with ER α and repression of its transcriptional activity at estrogen response elements (5,6). Recent studies have revealed that expression of <i>ESR2</i> is subject to epigenetic regulation and that loss of ER β expression positively contributes to epithelial-mesenchymal transition and enhanced invasiveness in prostate cancer (7,8). ER β has also been found to be negatively regulated at the posttranslational level through phosphorylation of its AF-1 domain, which promotes its ubiquitin-dependent proteasomal degradation (9,10).				
Background References		 Evans, R.M. (1988) Science 240, 889-95. Kuiper, G.G. et al. (1996) Proc Natl Acad Sci U S A 93, 5925-30. Moore, J.T. et al. (1998) Biochem Biophys Res Commun 247, 75-8. Leung, Y.K. et al. (2006) Proc Natl Acad Sci U S A 103, 13162-7. Hall, J.M. and McDonnell, D.P. (1999) Endocrinology 140, 5566-78. Pettersson, K. et al. (2000) Oncogene 19, 4970-8. Zhu, X. et al. (2004) Am J Pathol 164, 2003-12. Mak, P. et al. (2010) Cancer Cell 17, 319-32. Tateishi, Y. et al. (2006) Mol Cell Biol 26, 7966-76. Picard, N. et al. (2008) Mol Endocrinol 22, 317-30. 				

Species Reactivity Species reactiv

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key W: Western Blotting

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey

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